EFFECTS OF SOIL PROPERTIES ON COPPER SPECIATION IN SOIL SOLUTION

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Abstract

The numerical speciation analysis relies on quantitative assessment of concentrations of different forms of an element in soil solution (free ions, complex ions, neutral complexes), which have different abilities to react (ion activity). The reactivity affects the element's mobility and bioavailability. This method can be employed to estimate potential bioavailability and toxicity of a given element. This study was undertaken to evaluate effects of selected soil properties on changes in the total concentration of copper (Cu) and percentages of particular forms of this element in soil solution.

The study was based on a microplot experiment. The investigated factors were the soil texture, pH, organic carbon content and degree of soil copper contamination. Soil solutions were obtained with the vacuum displacement method. The concentration of copper in soil solution was determined with the ICP method, and the percentages of particular copper forms in the total copper concentration were calculated with the MINTEQA 2 software. It was found that copper in the analysed soil solutions occurred mainly in the form of metalorganic complexes. The increasing soil acidity was correlated with an increased percentage of free copper ions and copper complexes with organic matter. Simultaneously, the share of bonds with carbonates, sulfates and hydroxyl groups decreased. A decrease in the percentage of Cu^{2+} free ions in the soil solution was observed in response to an increasing organic carbon content, while the percentage of copper complexes with organic matter rose. The degree of soil copper contamination and soil texture had no influence on the percentages of different copper forms in the soil solution.

Key words: soil solution, copper, soil properties, numerical speciation, MINTEQA.

WPLYW WŁAŚCIWOŚCI GLEBY NA SPECJACJĘ MIEDZI W ROZTWORZE GLEBOWYM

Abstrakt

Numeryczna analiza specjacyjna polega na ilościowej ocenie stężenia różnych form badanego pierwiastka w roztworze glebowym (wolne jony, jony kompleksowe, obojętne kompleksy), które charakteryzują się różną zdolnością do reagowania (aktywnością jonów), co wpływa na ich ruchliwość i biodostępność. Metoda ta może więc być narzędziem umożliwiającym prognozowanie ich potencjalnej biodostępności lub toksyczności dla organizmów. W związku z tym podjęto badania dotyczące oceny wpływu wybranych właściwości gleby na zmiany całkowitego stężenia Cu oraz udziału poszczególnych form tego pierwiastka w roztworze glebowym.

W doświadczeniu mikropoletkowym badanymi czynnikami były skład granulometryczny, pH, zawartość Corg. w glebie oraz stopień zanieczyszczenia gleby miedzią. Roztwór glebowy pozyskiwano metodą podciśnieniową. Stężenie miedzi w roztworze glebowym oznaczono metodą ICP, a udział poszczególnych form miedzi w całkowitym stężeniu tego pierwiastka w roztworze glebowym obliczono za pomocą programu komputerowego MINTEQA 2. Stwierdzono, że w analizowanych roztworach glebowych miedź występuje głównie w formie kompleksów metaloorganicznych. Wraz ze wzrostem zakwaszenia zwiększał się w roztworze glebowym udział wolnych jonów i kompleksów miedzi z materią organiczną, a zmniejszał udział połączeń z węglanami, siarczanami i grupami hydroksylowymi. Wraz ze wzrostem zawartości C-org. w glebie, zmniejszał się w roztworze glebowym udział wolnych jonów Cu^{2+} , a zwiększał udział kompleksów miedzi z materią organiczną. Stopień zanieczyszczenia gleby metalami ciężkimi i skład granulometryczny nie wpływały na procentowy udział różnych form miedzi w roztworze glebowym.

Słowa kluczowe: roztwór glebowy, miedź, właściwości gleby, specjacja numeryczna, MINTEQA.

INTRODUCTION

The chemical composition of soil solution can be exploited to predict both bioavailability of trace elements to plants as well as potential contamination of plants grown for feed or food (HIRSH, BANIN 1990, KHOSHGOFTAR et al. 2004). Considering the number and diversity of chemical forms of trace elements in soil solution, special attention should be paid to the method of numerical speciation analysis. The method is based on quantifying concentrations of various forms of a given element in soil solution in an attempt to calculate its bioavailability or toxicity to living organisms (PARKER, PEDLER 1986, CHECKAI et al. 1987, MINNICH et al. 1987, CANCÉS et al. 2003).

The aim of the study was to determine the total concentration of copper (Cu) and percentages of individual chemical forms of this element in soil solution depending on such soil properties as the soil grain size composition, soil reaction, organic carbon content and degree of soil copper contamination.

MATERIAL AND METHODS

The study was based on a microplot experiment run at the Experimental Station of Agriculture, the Faculty of Biology at Warsaw University of Life Sciences in Skierniewice (central Poland). The microplots were represented by stoneware pots (1.2 m deep and 40 cm wide), which were filled with soil and placed outdoors.

The experimental factors were:

- three levels of soil pH: 4, 5 and 6;
- two soils: loamy sand (7% clay, 6% silt, 87% sand), sandy loam (13% clay, 5% silt, 82% sand);
- three levels of the content of soil organic carbon: 6, 9 and 12g C kg⁻¹;
- four degrees of soil contamination with Cu: natural content – 0 (10 mg kg⁻¹), I (30 mg kg⁻¹), II (50 mg kg⁻¹) and III (80 mg kg⁻¹) (KABATA-PENDIAS et al. 1993).

The experiment consisted of 216 microplots, which represented 72 combinations of the investigated factors. All the factors were observed in three replications arranged in a split-plot experimental design. The experimental samples were collected from each microplot's topmost soil layer (0-30 cm down). The soil samples were dried up to the level of air humidity. Afterwards, they were ground in a china mortar and put through a sieve with the 1 mm diameter mesh net. The samples thus prepared were used to obtain soil solutions according to the vacuum displacement method with a vacuum pump (Dynavac OP4). The total concentration of copper was determined by Inductively Coupled Plasma Atomic Emission Spectrometry (ICP-AES method) (apparatus: IRYS Advantage ThermoElementar). The MINTEQA2 for Windows software was used for calculations of shares of particular forms of Cu ions in soil solution. The results were statistically analyzed with ANOVA and simple linear regression. The differences between means were analysed with the Tukey's test at $p=0.05$. Statistical analyses were conducted with Statgraphics 5.1. software.

RESEARCH RESULTS

The total concentration of Cu in the analysed soil solutions ranged from 1.23 to 2.25 $\mu\text{mol dm}^{-3}$ and differed depending on soil properties.

The results of ANOVA showed that the total concentration of copper in soil solutions increased significantly as the soil reaction became more acidic and the degree of copper soil contamination increased. The concentration of Cu was higher in soil solution of loamy sand than of sandy loam. The content of organic carbon did not significantly affect Cu concentrations in soil solution (Table 1).

Table 1

Total Cu concentration ($\mu\text{mol dm}^{-3}$) versus soil properties

Item	Soil texture		Soil reaction (pH_{KCl})			Content of organic carbon (g kg^{-1})			Degree of Cu soil contamination			
	loamy sand	sandy loam	4	5	6	6.0	9.0	12.0	0	I	II	III
Cu	1.76	1.26	2.05	1.46	1.38	1.52	1.52	1.49	1.23	1.31	1.75	2.25
$\text{LSD}_{0.05}$	0.18		0.21			0.27			0.25			

The results of speciation analysis of copper in soil solutions indicated the highest percentage of Cu complexes with organic matter and carbonates. The prevalent forms of copper were Cu-org. complexes, whose average share ranged from 28.2 to 93.2% of the total concentration of copper in soil solution. The second most abundant form of copper observed in soil solutions were Cu-CO₃ complexes, whose contribution to total copper ranged from 12.7 to 62.3%, depending on soil properties (Tables 2–5).

Table 2

Percentages of different copper forms in soil solution according to soil pH value (pH_{KCl})

Form of Cu	Soil pH		
	4.0	5.0	6.0
Cu^{2+}	6.5	5.9	4.4
Cu-org.	93.2	56.2	28.2
CuOH^+	tc*	3.0	3.4
Cu-CO_3	tc	34.6	62.3
$\text{Cu}(\text{CO}_3)_2^{2-}$	tc	tc	1.4

*trace content

The highest percentage (93.2%) of Cu-org. complexes in the total concentration of copper was observed in the soil solutions of most acid soils (Table 2). As the soil pH was increasing, amounts of metalorganic complexes of copper were steadily decreasing, while the percentage of Cu-CO₃ complexes was growing. Copper complexes with carbonates reached 34.6% in the solutions of soils with pH 5.0 and 62.3% in the solutions of soils with pH 6.0. Other forms of copper (CuOH^+ , Cu-CO₃ and $\text{Cu}(\text{CO}_3)_2^{2-}$) were not observed in the solutions of soils with the lowest pH value. At the same time, their share in the total Cu concentration in solutions of soils with higher pH was relatively low (Table 2).

Regardless of the degree of soil Cu contamination, the dominant forms of copper in soil solution were Cu-org. complexes, which approximately made up 59% of this element's total soil solution concentration. At the same time, a relatively high percentage (32%) of Cu-CO₃ complexes was observed. In both Cu unpolluted and contaminated soils, an average share of free ions Cu²⁺ was just 5.5% of the total concentration of copper in soil solution (Table 3).

Table 3

Percentages of different copper forms in soil solutions according to soil copper contamination

Form of Cu	Degree of Cu soil contamination			
	0	I	II	III
Cu ²⁺	5.4	5.4	5.7	5.8
Cu-org.	59.4	58.6	59.0	59.8
CuOH ⁺	2.1	2.1	2.2	2.1
Cu-CO ₃	33.0	33.0	32.3	31.6
Cu(CO ₃) ₂ ²⁻	0.5	0.5	0.5	0.5

Table 4

Percentages of different copper forms in soil solution according to content of soil organic carbon

Form of Cu	Content of soil organic carbon (g kg ⁻¹)		
	6.0	9.0	12.0
Cu ²⁺	6.5	5.3	5.0
Cu-org.	37.1	61.7	78.8
CuOH ⁺	2.6	2.1	1.8
Cu-CO ₃	52.6	30.1	14.2
Cu(CO ₃) ₂ ²⁻	0.9	0.5	-

The effect of the content of soil organic carbon on copper speciation in soil solutions was considerable. A two-fold increase of soil organic carbon content nearly doubled the share of Cu-org. in the total concentration of copper in soil solution (from 37.1% to 78.8%). Simultaneously, the percentage of Cu-CO₃ complexes decreased three-fold (from 52.6% to 14.2%). Free ions Cu²⁺ made up 5.0% of all Cu forms in soil solutions of the soils characterized by the highest content of organic carbon, and 6.5% in soil solution of soils with the lowest content of organic carbon. Other forms of copper represented low percentages in the total content of copper in the soil solution (Table 4).

Soil texture had a weak effect on the share of copper forms in the distribution of this element in soil solution. The Cu-org. form was prevalent in the solutions of all the analysed soils: an average share of this form in the total copper concentration in soil solutions was 56.7% in sandy loam soil and 60.8% in loamy sand. The Cu-CO₃ complexes had a large share in the overall distribution of copper forms in soil solutions. Unlike Cu-org., they were more abundant in sandy loam soil solutions (on average 34.5%). In loamy sand soils, complexes of copper with carbonates constituted 30.0% of the total amount of copper in the soil solution (Table 5).

Table 5

Percentages of different copper forms according
to soil texture

Form of Cu	Soil texture	
	loamy sand	sandy loam
Cu ²⁺	5.8	5.6
Cu-org.	60.8	56.7
CuOH ⁺	2.9	2.3
Cu-CO ₃	30.0	34.5
Cu(CO ₃) ₂ ²⁻	0.5	0.5

DISCUSSION

The results of the present study suggest a significant effect of soil properties on the total concentration of copper in soil solution. In solutions of the analysed soils, total copper concentrations increased due to soil Cu contamination and in response to soil acidification. At the same time, the copper total concentration was higher in solution of loamy sand soil. On the other hand, no effect of the soil organic carbon content on the total concentration of Cu in soil solution was observed (Table 1). Such influence of soil properties on copper concentration in soil solution has been confirmed by other authors (JEFFERY, UREN 1983, McGRATH et al. 1988, SAUVÉ et al. 1997). The observed effects are attributed to the high affinity of copper for organic matter (NORVELL 1991).

The observed high ratio of Cu-org. complexes in the total concentration of copper in soil solution is confirmed in literature (SANDERS 1982, 1983, McBRIDE, BOULDIN 1984, FOTOVAT, NAIDU 1997). The share of metalorganic complexes estimated in this experiment at 28.2 to 93.2% corresponds well with the results of SMAL (1999), which were obtained with the use of SOILCHEM

software. The latter showed that copper organic complexes amounted to 75.4% of the total concentration of this element in soil solution of a humus layer of forest soils and 48.4% in agricultural soils. The results of the numerical analysis of organic soil solutions carried out by RUTKOWSKA (2008) with the MINTEQA2 software indicated that the percentage share of Cu-org. complexes in the total concentration of copper in soil solution ranged from 74 to 86%. Even a higher percentage of copper organic complexes, reaching 99.5%, was found by McBRIDE and BOULDIN (1984).

In the analysed solutions, copper was also found in the form of complexes with carbonates, whose share ranged from 12.7 to 62.3% of the total concentration of copper in soil solution, depending on soil reaction and soil richness in organic carbon (Tables 2-5). The share of Cu-CO₃ complexes in soil solution decreased with an increase of soil organic carbon content and a change of soil reaction into acid one. SMAL (1999), SAEKI et al. (2002) and RUTKOWSKA (2008) confirmed presence of copper in the form of Cu-CO₃ complexes in soil solution. However, none of these authors obtained a percentage of this form in soil solution as high as observed in the present study. RUTKOWSKA (2008) showed that solutions of agricultural soils contain complexes of copper with carbonates on a level ranging from 5.0 to 6.8% of the total concentration of Cu in soil solution.

Regardless of the soil properties investigated, a small percentage of free Cu²⁺ ions was observed in the total concentration of copper in soil solution. Similar observations were reported by NOLAN et al. (2003) who showed a 5% share of free Cu²⁺ ions in the soil solutions analysed in their research. A small share of free Cu²⁺ in soil solution, no more than slightly over 12%, was also observed by SANDERS (1983), McGRATH et al. (1988) and FOTOVAT, NAIDU (1997). According to YUAN (2009), WÓJCIKOWSKA-KAPUSTA, NIEMCZUK (2009) and JAWORSKA, DĄBKOWSKA-NASKRĘT (2012) this result is due to the strong binding of copper by the soil solid phase. As claimed by CANCEŚ et al. (2003), speciation of copper in soil solution is conditioned by soluble forms of organic substance. SALAM, HELMKE (1998) suggest that complexing is a reaction that substantially conditions the course of copper speciation in soil solution.

According to SMAL (1999), SAEKI et al. (2002), ARTIOLA (2005) and RUTKOWSKA (2008) copper in soil solution can also form bonds with other nonorganic ligands (e.g. SO₄²⁻, NH₃, NO₃⁻, H₂PO₄²⁻, Cl⁻, OH⁻). In the numerical speciation analysis carried out in this study, a small proportion of CuOH⁺ bonds was detected in the analysed soil solutions, but other forms were not observed (Tables 2-5).

CONCLUSIONS

Based on the numerical speciation analysis of copper in the solutions of soils with different physical and chemical properties it was concluded that:

- the estimates derived by running a MINTEQA2 software package of the percentages of ion forms in soil solutions showed that copper occurred in soil solution in the form of complexes with humus acids;
- among the soil properties studied, soil reaction had the strongest influence on the per cent shares of copper forms in soil solution; in a more acid soil, the ratio of copper free ions and copper complexes with organic matter, bonds with carbonates as well as sulfates and hydroxyl groups decreased;
- the degree of soil Cu contamination and soil texture had no effect on percentages of different forms of copper in soil solution;
- with a higher organic carbon content in soil, soil solution contained a lower share of free copper ions, whereas the share of copper complexes with organic matter increased.

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